

| Project Title | Funding | Strategic Plan Objective | Institution |
|--|-------------|--------------------------|---|
| Vasopressin receptors and social attachment | \$121,500 | Q4.S.B | Emory University |
| Using zebrafish and chemical screening to define function of autism genes | \$399,999 | Q4.S.B | Whitehead Institute for Biomedical Research |
| Using iPS cells to study genetically defined forms with autism | \$200,000 | Q4.S.B | Stanford University |
| Using induced pluripotent stem cells to identify cellular phenotypes of autism | \$800,000 | Q4.S.B | Stanford University |
| Using Drosophila to model the synaptic function of the autism-linked NHE9 | \$150,000 | Q4.S.B | Massachusetts Institute of Technology |
| Transgenic mouse model to address heterogeneity in autism spectrum disorders | \$468,586 | Q4.S.B | Vanderbilt University |
| The role of SHANK3 in the etiology of autism spectrum disorder | \$28,000 | Q4.S.B | Johns Hopkins University |
| The role of SHANK3 in autism spectrum disorders | \$360,000 | Q4.S.B | Mount Sinai School of Medicine |
| The genetics of restricted, repetitive behavior: An inbred mouse model | \$60,000 | Q4.S.B | University of Florida |
| The genetic control of social behavior in the mouse | \$346,000 | Q4.S.B | University of Hawai'i at Manoa |
| Systematic analysis of neural circuitry in mouse models of autism | \$149,973 | Q4.S.B | Cold Spring Harbor Laboratory |
| Synaptic plasticity, memory and social behavior | \$52,154 | Q4.S.B | New York University |
| Synaptic deficits of iPS cell-derived neurons from patients with autism | \$86,588 | Q4.S.B | Stanford University |
| Synaptic and circuitry mechanisms of repetitive behaviors in autism | \$400,000 | Q4.S.B | Massachusetts Institute of Technology |
| Small-molecule compounds for treating autism spectrum disorders | \$175,000 | Q4.S.B | The University of North Carolina at Chapel Hill |
| Shank3 mutant characterization in vivo | \$28,000 | Q4.S.B | University of Texas Southwestern Medical Center |
| Serotonin, corpus callosum, and autism | \$303,250 | Q4.S.B | University of Mississippi Medical Center |
| Serotonin, autism, and investigating cell types for CNS disorders | \$90,000 | Q4.S.B | The Rockefeller University |
| Role of UBE3A in neocortical plasticity and function | \$490,000 | Q4.S.B | Duke University |
| Role of L-type calcium channels in hippocampal neuronal network activity | \$32,741 | Q4.S.B | Stanford University |
| Role of a novel Wnt pathway in autism spectrum disorders | \$750,000 | Q4.S.B | University of California, San Francisco |
| Regulation of gene expression in the brain | \$2,086,763 | Q4.S.B | National Institutes of Health |
| Preclinical testing of novel oxytocin receptor activators in models of autism phenotypes | \$167,572 | Q4.S.B | University of North Carolina at Chapel Hill |
| Preclinical testing of novel oxytocin receptor activators in models of autism phenotypes | \$39,325 | Q4.S.B | University of North Carolina at Chapel Hill |
| Preclinical testing of novel oxytocin receptor activators in models of autism phenotypes | \$346,289 | Q4.S.B | University of North Carolina at Chapel Hill |
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| Patient iPS cells with copy number variations to model neuropsychiatric disorders | \$207,388 | Q4.S.B | The Hospital for Sick Children |
| Optimization of methods for production of both ICSI- and SCNT derived baboon embryonic stem cells | \$260,102 | Q4.S.B | Southwest Foundation For Biomedical Research |
| NrCAM, a candidate susceptibility gene for visual processing deficits in autism | \$0 | Q4.S.B | University of North Carolina at Chapel Hill |
| Novel strategies to manipulate Ube3a expression for the treatment of autism and Angelman syndrome | \$0 | Q4.S.B | University of North Carolina at Chapel Hill |
| Novel probiotic therapies for autism | \$570,145 | Q4.S.B | California Institute of Technology |
| Novel models to define the genetic basis of autism | \$289,633 | Q4.S.B | Cold Spring Harbor Laboratory |
| Novel genetic animal models of autism | \$274,750 | Q4.S.B | University of Texas Southwestern Medical Center |
| Novel, subtype selective potentiators of nicotinic acetylcholine receptors | \$325,757 | Q4.Other | University of Alaska Fairbanks |
| Neuropharmacology of motivation and reinforcement in mouse models of autistic spectrum disorders | \$0 | Q4.S.B | University of North Carolina School of Medicine |
| Neurexin function in vivo: Implications for autism and mental retardation | \$392,500 | Q4.S.B | University of Texas Southwestern Medical Center |
| Neurogenetics in a model for procedural learning | \$33,053 | Q4.S.B | University of California, Los Angeles |
| Neurogenetic model of social behavior heterogeneity in autism spectrum disorders | \$795,188 | Q4.S.B | Duke University |
| Neurobiology of sociability in a mouse model system relevant to autism | \$354,375 | Q4.S.B | University of Pennsylvania |
| Neurobiology of mouse models for human chr 16p11.2 microdeletion and fragile X | \$210,000 | Q4.S.B | Massachusetts Institute of Technology |
| Neurobiological mechanism of 15q11-13 duplication autism spectrum disorder | \$304,500 | Q4.S.B | Beth Israel Deaconess Medical Center |
| Neural mechanisms of social cognition and bonding | \$0 | Q4.S.B | Emory University |
| Neural mechanisms of social cognition and bonding | \$43,862 | Q4.S.B | Emory University |
| Neural and cognitive mechanisms of autism | \$375,000 | Q4.S.B | Massachusetts Institute of Technology |
| Murine genetic models of autism | \$172,389 | Q4.S.B | Vanderbilt University |
| Mouse genetic model of a dysregulated serotonin transporter variant associated with autism | \$0 | Q4.S.B | Vanderbilt University |
| Modeling and pharmacologic treatment of autism spectrum disorders in Drosophila | \$127,500 | Q4.S.B | Albert Einstein College of Medicine of Yeshiva University |
| Micro-RNA regulation in pluripotent stem cells | \$19,189 | Q4.S.B | Southwest Foundation For Biomedical Research |
| Mice lacking Shank postsynaptic scaffolds as an animal model of autism | \$128,445 | Q4.S.B | Massachusetts Institute of Technology |
| Methods for production of ICSI and SCNT derived macaque stem cells | \$19,188 | Q4.S.B | Southwest Foundation For Biomedical Research |
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| Investigation of the role of MET kinase in autism | \$366,308 | Q4.S.B | Johns Hopkins University School of Medicine |
| Investigating the effects of chromosome 22q11.2 deletions | \$150,000 | Q4.S.B | Columbia University |
| Interaction between MEF2 and MECP2 in the pathogenesis of autism spectrum disorders -2 | \$0 | Q4.S.B | Burnham Institute |
| Interaction between MEF2 and MECP2 in the pathogenesis of autism spectrum disorders - 1 | \$0 | Q4.S.B | Burnham Institute |
| Integrated approach to the neurobiology of autism spectrum disorders | \$232,118 | Q4.S.B | Yale University |
| Insight into MeCP2 function raises therapeutic possibilities for Rett syndrome | \$295,298 | Q4.S.B | University of California, San Francisco |
| Identifying impairments in synaptic connectivity in mouse models of ASD | \$40,000 | Q4.S.B | University of Texas Southwestern Medical Center |
| Identifying genetic modifiers of rett syndrome in the mouse | \$30,000 | Q4.S.B | Baylor College of Medicine |
| Identification of autism genes that regulate synaptic Nrx/Nlg signaling complexes | \$200,000 | Q4.S.B | Stanford University |
| High-resolution diffusion tensor imaging in mouse models relevant to autism | \$199,724 | Q4.S.B | University of Pennsylvania |
| High content screens of neuronal development for autism research | \$210,977 | Q4.S.B | University of California, San Diego |
| Genomic resources for identifying genes regulating social behavior | \$60,000 | Q4.S.B | Emory University |
| Genomic imbalances at the 22q11 locus and predisposition to autism | \$400,000 | Q4.S.B | Columbia University |
| Genetic models of serotonin transporter regulation linked to mental disorders | \$184,375 | Q4.S.B | Medical University of South Carolina |
| Function and dysfunction of neuroligins | \$374,383 | Q4.S.B | Stanford University |
| Functional study of synaptic scaffold protein SHANK3 and autism mouse model | \$150,000 | Q4.S.B | Duke University |
| Functional genomic dissection of language-related disorders | \$235,753 | Q4.S.B | University of Oxford |
| Exploring the neuronal phenotype of autism spectrum disorders using induced pluripotent stem cells | \$241,503 | Q4.S.B | Stanford University |
| Evaluation of altered fatty acid metabolism via gas chromatography/mass spectroscopy and time-of-flight secondary ion mass spectroscopy imaging in the propionic acid rat model of autism spectrum disorders | \$25,000 | Q4.S.B | University of Western Ontario |
| Dynamic regulation of Shank3 and ASD | \$300,000 | Q4.S.B | Johns Hopkins University |
| Dissecting the neural control of social attachment | \$772,500 | Q4.S.B | University of California, San Francisco |
| Dissecting the circuitry basis of autistic-like behaviors in mice | \$175,000 | Q4.S.B | Massachusetts Institute of Technology |

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| Development of a high-content neuronal assay to screen therapeutics for the treatment of cognitive dysfunction in autism spectrum disorders | \$0 | Q4.S.B | Massachusetts Institute of Technology |
| Design & synthesis of novel CNS-active oxytocin and vasopressin receptor ligands | \$560,535 | Q4.Other | The Scripps Research Institute |
| Deriving neuroprogenitor cells from peripheral blood of individuals with autism | \$0 | Q4.S.B | University of Utah |
| Control of synaptic protein synthesis in the pathogenesis and therapy of autism | \$155,063 | Q4.S.B | Massachusetts General Hospital |
| Cntnap2 in a behavioral model of autism | \$262,356 | Q4.S.B | University of California, Los Angeles |
| Characterization of the transcriptome in an emerging model for social behavior | \$426,250 | Q4.S.B | Emory University |
| Characterization of autism susceptibility genes on chromosome 15q11-13 | \$47,606 | Q4.S.B | Beth Israel Deaconess Medical Center |
| Characterization of a novel mouse model of restricted repetitive behaviors | \$222,000 | Q4.S.B | University of North Carolina at Chapel Hill |
| Central vasopressin receptors and affiliation | \$32,896 | Q4.S.B | Emory University |
| Central vasopressin receptors and affiliation | \$364,425 | Q4.S.B | Emory University |
| Cellular and genetic correlates of increased head size in autism spectrum disorder | \$282,901 | Q4.S.B | Yale University |
| Caspr2 dysfunction in autism spectrum disorders | \$0 | Q4.S.B | Yale University |
| Behavioral and physiological consequences of disrupted Met signaling | \$800,000 | Q4.S.B | University of Southern California |
| Basal ganglia circuitry and molecules in pathogenesis of motor stereotypy | \$387,767 | Q4.S.B | University of California, Los Angeles |
| Autism iPSCs for studying function and dysfunction in human neural development | \$254,152 | Q4.S.B | The Scripps Research Institute |
| A preclinical model for determining the role of AVPR1A in autism spectrum disorders | \$0 | Q4.S.B | Mount Sinai School of Medicine |
| A novel cell-based assay for autism research and drug discovery | \$60,000 | Q4.S.B | University of Arizona |
| Animal models of neuropsychiatric disorders | \$1,769,941 | Q4.S.B | National Institutes of Health |
| Animal models of autism: Pathogenesis and treatment | \$84,999 | Q4.S.B | University of Texas Southwestern Medical Center |
| Animal model of speech sound processing in autism | \$325,125 | Q4.S.B | University of Texas at Dallas |
| Analysis of cortical circuits related to ASD gene candidates | \$0 | Q4.S.B | Cold Spring Harbor Laboratory |
| A mouse knock-in model for ENGRAILED 2 autism susceptibility | \$227,135 | Q4.S.B | University of Medicine & Dentistry of New Jersey |
| 16p11.2: defining the gene(s) responsible | \$175,000 | Q4.S.B | Cold Spring Harbor Laboratory |

